

**K802391 EMG-UT 40**Jan 9, 1981  
99 days to decisionK802391 · Product code: **HCC** · Neurology  
Source: <https://www.510kdatabase.net/k802391/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Oct 2, 1980
Decision date	Jan 9, 1981
Days to decision	99 days
Third-party review	No

**APPLICANT**

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Company	<b>American Scientific Corp.</b>
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k802391/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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